I. CLAIMS

1. (Currently Amended) A tablet for oral administration, which disintegrates in the oral cavity within 60 seconds, consisting essentially of (i) a therapeutically effective amount of an active ingredient, (ii) an effective amount of spray-dried mannitol as primary disintegrant, of which at least 80% has an average particle size over 100 um, (iii) an effective amount of crospovidone as secondary disintegrant, and (iv) one or more pharmaceutically acceptable excipients, the tablet containing no microcrystalline cellulose.

2. (Original) The tablet of claim 1, wherein the contents of the spray-dried mannitol and the crospovidone are in the ranges of 30 to 95% and 1 to 10% by weight, respectively, based on total weight of the tablet.

3. (Original) The tablet of claim 1, wherein the active ingredient is selected front the group consisting of acetominophen, domperidone, famotidine, meclizine hydrochloride, scopolamine hydrobromide, ondansetron HCl, cisapride, granisetron, sildenafil, loratadine and amlodipine.

4. (Original): A process for the preparation of a tablet for oral administration which disintegrates in the oral cavity within 60 seconds, comprising direct-compressing a mixture consisting essentially of (i) a therapeutically effective amount of an active ingredient, (ii) spray-dried mannitol, (iii) crospovidone, and (iv) one or more pharmaceutically acceptable water soluble excipients other than microcrystalline cellulose.